Your Ref.: *NSC-11.926* 



11 Publication number:

(12)

# **EUROPEAN PATENT APPLICATION**

(21) Application number: 87104765.0

(5) Int. Cl.4: **F01N 3/28**, B01D 53/36

2 Date of filing: 31.03.87

Priority: 31.03.86 JP 71273/86

43 Date of publication of application: 04.11.87 Bulletin 87/45

Designated Contracting States: DE

1 Applicant: NIPPON STEEL CORPORATION 6-3 Otemachi 2-chome Chiyoda-ku Tokyo 100(JP)

Applicant: NIPPON KINZOKU CO., LTD. 6-18, 3-chome, Kamiya Kita-ku Tokyo(JP)

Inventor: Nakamura, Yasushi c/o Nippon Steel

R & D Laboratories-I 1618, Ida, Nakahara-ku Kawasaki-shi, Kanagawa (JP)

Inventor: Yamanaka, Mikio c/o Nippon Steel

R & D Laboratories-II 5-10-1, Fuchinobe

Sagamihara-shi Kanagawa(JP)

Inventor: Oomura, Keiichi c/o Nippon Steel

Corp.

R & D Laboratories-II 5-10-1, Fuchinobe

Sagamihara-shi Kanagawa(JP) Inventor: Arakawa, Motohiko

c/o Nippon Steel Corp.6-3 Otemachi

2-chome

Chiyoda-ku Tokyo(JP)

Inventor: Yashiro, Toshiyuki

c/o NIKKIN R & D Co., LTD. 4-10-1, Funado

Itabashi-ku Tokyo(JP) Inventor: Izumi, Masaki

c/o Nippon Kinzoku Co., Ltd. 6-18, Kamiya

3-chome

Kita-ku Tokyo(JP)

(74) Representative: Kador & Partner Corneliusstrasse 15 D-8000 München 5(DE)

Method for producing a base of a catalyst carrier for automobile exhaust gas-purification.

 $\bigcirc$  In the carrier of a catalyst for purifying the exhaust gas, the  $\gamma$ -Al<sub>2</sub>O<sub>2</sub> powder, on which the catalyst is impregnated, is carried on an Al<sub>2</sub>O<sub>2</sub> film formed on a stainless steel foil formed into a honeycomb. The honeycomb is formed by coiling a corrugated sheet and a flat sheet together, and is conventionally bonded with an outer cylinder on the entire peripheral surface thereof by an Ni-brazing material. In the present invention, an Al (alloy)-clad foil or hot-dip Al (alloy) galvanized foil is used and heat treated to simultaneously bond the foils together and the honeycomb to the outer cylinder, to thereby form the Al<sub>2</sub>O<sub>3</sub> film, and to diffuse Al into the

stainless steel foil.

### 0 243 702

# $\frac{\text{METHOD FOR PRODUCING A BASE OF A CATALYST CARRIER FOR AUTOMOBILE EXHAUST GAS-PURIFICATION}{\text{PURIFICATION}}$

# BACKGROUND OF THE INVENTION -- -

#### I. Field of the Invention

5

The present invention relates to a method for producing a base of a catalyst carrier used for the exhaust gas purification device of an automobile.

#### 2. Description of the Related Art

Since nineteen-seventies, air pollution by automobile emission control has come into a social problem, and regulations for the automobile exhaust emissions have been promulgated to the effect that passenger automobiles must be equipped with a device for purifying the exhaust gas. Several systems have been proposed for devices for purifying the exhaust gas, but the most widely used at present is a catalyst converter system whereby HC and CO are oxidized and, simultaneously, NOx is reduced. These catalyst converters is made up of a ceramic honeycomb of corderite, mounted in a metallic cylinder, as a fundamental body, on which porous  $\gamma$ -Al<sub>2</sub>O<sub>3</sub> powder, impregnated with a noble metal (Pt  $\sim$  Pd) catalyst, is deposited. The ceramic honeycomb, however, has disadvantages in that it is not highly resistant to mechanical impact, and in addition, the exhaust resistance is undesirably high. The techniques disclosed to cope with these drawbacks include a known metal honeycomb in which metal foils of several tens  $\mu$ m in the form of a flat sheet and corrugated one are coiled into a roll inserted mounted in a outer case of stainless steel.

Japanese Examined Patent Publication No. 57-55,886, discloses an improvement of the above method by a technique in which the metal honeycomb is fixed to the outer case at the ends of body by electron beam welding or by brazing, or improved by the technique disclosed in Japanese Examined Patent Publication No. 58-23,I38. These techniques, however, have various drawbacks. For example, a high Alcontaining stainless steel used for the foil because of its excellent oxidation resistance has a poor rollability and is expensive. Also, from the viewpoint of the rollability, limitations should be imposed on the amount of alloying elements, which makes it difficult to attain a stable formation of an Al<sub>2</sub>O<sub>2</sub> film necessary for maintaining a sufficient oxidation resistance for a long time. In attempts to eliminate these drawbacks, Japanese Unexamined Patent Publication No. 54-97,593, Japanese Examined Patent Publication No. 57-3,418, Japanese Unexamined Patent Publication No. 54-33,888, and German Patent No. 2.745,188 disclose to give iron, stainless steel, heat-resistant alloy or the like, an Al coating, and then carry out a chemical treatment or heat treatment in an oxidizing atmosphere, thereby forming an Al<sub>2</sub>O<sub>3</sub> film or alloy.

In the manufacture of a metal substrate, the metal foils are formed into a honeycomb, followed by the bonding process between corrugated sheets and flat sheets as well as bonding between the outer cylinder and honeycomb. Generally, this bonding is carried out by brazing. The brazing material used for this bonding must be a high Ni-brazing material with a high melting point and a high heat resistance, considering the severe condition under the usage of the brazed assembly. Thus, Ni is oxidized at the surface of brazed parts by the exhaust gas. Since the Ni oxide has an effect of catalysis on unburnt components in the exhaust gas, a reaction takes place at the brazed parts, causing a partial raise in temperature at these parts. Accordingly, the brazed parts may melt down, if the Ni-brazing material has a melting point of less than Il00°C. Accordingly, it is necessary to use a brazing material having a melting point of Il00°C or more. Therefore, the heat treatment must be carried out at a temperature of Il00°C or more, in vacuum, for a perfect brazing.

In addition, because of a high Al content of the substrate, the wettability and the flowability of the brazing material are greatly reduced on the substrate surface. Accordingly, the high temperature necessary for the heat treatment lowers the production efficiency and increases costs. Furthermore, the brazed parts of a honeycomb manufactured as described above contain only a small amount of Al diffused from the substrate material, with the result that an  $Al_2O_2$  film having an excellent heat-resistance cannot be formed. Thus, the heat-resistance of the brazed parts is lower than that of the substrate material and the adhesion of  $\gamma$ -Al<sub>2</sub>O<sub>3</sub> powder is poor.

The catalyst carrier is exposed to severe operating conditions, for example..a high speed exhaust gas flow and extreme mechanical vibration upon acceleration. Therefore, the bonding strength of a carrier is an another important factor in the durability of the device. As is clear from the above explanation, to enable the use of Al-coated material for the substrate of a catalyst carrier, a novel bonding technique must be developed.

#### SUMMARY OF THE INVENTION

10

35

The present invention is intended to eliminate the above disadvantages.

Therefore, according to the present invention, the mutual bonding of parts of a honeycomb and the bonding between the honeycomb and the outer cylinder by using the Al or Al-alloy coating as the brazing material, thus eliminating the need for an additional brazing material, while imparting the very high Al content to the substrate required for the formation of a protective Al<sub>2</sub>O<sub>3</sub> film and a high heat resistance without trouble in cold rolling of the substrate foil.

Namely, the present invention is related to a method for producing a base of a catalyst carrier for a device for the purification of the exhaust gas of an automobile, and is characterized by rolling a stainless-steel sheet having a layer consisting of Al or Al-alloy clad on one or both surfaces of the stainless-steel sheet to form a composite (clad) foil, producing a honeycomb structure by using the composite foil, mounting the honeycomb structure in an outer cylinder of heat-resistant steel, to form a carrier base, and heat-treating the carrier base in vacuum or a reducing atmosphere at a temperature of 600 to  $1300^{\circ}$ C. In the present invention, the stainless steel strip is the core material of the composite foil, and the heat-resistance can be further enhanced when this core material is an Al-containing stainless steel. The stainless steel strip having a foil layer clad on one or both surfaces thereof may be a so-called "clad foil" which is obtained by rolling an Al (alloy) - stainless steel - Al (alloy) composite sheet, or may be an Al plated stainless-steel sheet by hot-dipping to a predetermined thickness. The reducing atmosphere mentioned above may conventionally contain hydrogen gas or the like, and in addition, may be produced by a reducing agent dispersed on the honeycomb. The base formed by the method according to the present invention is subsequently heated, if necessary, at a temperature of from 850 to  $1200^{\circ}$ C in air atmosphere, to form an  $Al_2O_3$  film on which  $\gamma$ -Al $_2O_3$  as the direct carrier of the catalyst is then deposited.

## DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention is now explained in detail.

The Al layer used in the present invention may be a layer of pure Al, and in addition, may consist of corrosion-resistant Al alloys or high-strength Al alloys containing Mn, Mg, Si, Cu, and the like.

The surface of the stainless steel strip used in the present invention is alloyed with the Al layer by the subsequent heat treatment, so that an improved heat-resistance is exhibited. Accordingly, any of the martensitic, austenitic, and dual-phase stainless steels can be used, provided that these steels contain 9% or more of Cr and are rollable. The desirable composition-series according to the present invention are 0.5% or less of C, 0  $\sim$  60% of Ni, 9  $\sim$  30% of Cr, 0  $\sim$  10% of one or more of Mo, W, and Co. 0  $\sim$  3% of one or more of Ti, Nb, Zr, and Hf, 0  $\sim$  0.2% of rare earth metal (REM) and Y. 0  $\sim$  0.01% of one or more of Mg and Ca, and , if necessary, a small amount of Cu and B; the balance being Fe and unavoidable impurities. Even if Al is not added to the stainless steel strip, the strip is enriched with Al by the subsequent heat treatment from the surface Al or Al alloy layer. Nevertheless, the heat-resistance of honeycomb is further enhanced if 6% or less of Al is preliminarily added to the stainless steel strip.

The stainless steel foil having foil layers of Al or Al-alloy clad on both surfaces thereof is then produced according to the following procedures. A material clad with Al or an Al-alloy is produced, or a hot-dip Al plating is carried out, in a process wherein the stainless steel strip has a thickness of from approximately 0.1 to 1.0 mm. The Al or Al-alloy coated or clad strip is further rolled until a foil thickness, e.g., from 30, to 70 µm, appropriate for the metal honeycomb is obtained. The clad material may be produced by any appropriate known method, e.g., a rolling method. In the hot-dip Al-plating, the formation of a Fe-Al alloy layer between Al and the substrate should be suppressed as thin as possible, to ensure an excellent rollability thereof after the plating. To this end, a small amount of Si should be added to the molten Al bath or a small amount of N preliminarily should be added to the stainless steel. The thickness ratio of the stainless-steel sheet to the Al or Al alloy layer is desirable from approximately 15:1 to approximately 3:1, at the time of forming the honeycomb. This thickness ratio determines the Al content, i.e., the heat-resistance

# 0 243 702

of a honeycomb after being subjected to a subsequent heat treatment. Generally, the heat-resistance is enhanced with an increase in the Al content, but an excessive Al content leads not only to an embrittlement of material but also to a lowering of the melting point, with the result that the catalyst carrier is not durable at a high temperature. The Al content is, therefore, preferably 30% by weight or less after the alloy is formed by cladding followed by heat treatment.

A composite foil produced as described above is subjected to corrugation shaping and is then overlapped with a composite foil in the form of a flat sheet. Both foils are then coiled in the form of a roll or is laminated to provide a honeycomb body, which is then press-mounted in an outer cylinder made of heat-resistant steel. The heat-resistant steel herein may be a heat-resistant casting, low-alloy steel, or stainless steel, provided that a satisfactory heat resistance of the container of the catalyst for the exhaust gas purification device is provided.

The honeycomb or the outer cylinder having the honeycomb mounted therein is heat-treated at a temperature of from 600 to I300°C in vacuum or a reducing atmosphere. The AI or AI-alloy present on the surface(s) of the honeycomb is then alloyed with the inner stainless steel by a mutual diffusion, and as s result, an excellent heat-resistance is given, while the contacting parts between the corrugated sheet and the flat sheet, as well as the contacting parts between the outermost peripheral surface of the honeycomb and the inner surface of the outer cylinder are fusion-adhered together by the high Al layer on the surface with a relatively low melting temperature, thereby providing a strong fixing therebetween. It is possible to enlarge the bonding area and smooth the bonding by inserting an Al foil or an Al-alloy foil between the outermost peripheral surface of the honeycomb and the inner surface of the outer cylinder. The vacuum and reducing atmosphere referred to herein, which is indispensable for the bonding of a honeycomb, is generally such that an oxide film having a thickness detrimental to the bonding is not formed on the surfaces of the high Al layer during the heat treatment for bonding. Specifically, the vacuum must have pressure of 10<sup>-4</sup> Torr or less, and the reducing atmosphere must be non-oxidizing gas, e.g., dry hydrogen. or dry nitrogen, argon, helium or the like, with a hydrogen content of 5% or more. To create the reducing atmosphere necessary for the above bonding, Al, Ti, Zn, or polyethyleneimine in powder or liquid form, which are active elements or compounds, is coated on the surfaces of honeycomb, and the heat treatment is then carried out in an inert atmosphere of argon gas or the like, or in an air atmosphere while sealing the honeycomb with another foil of stainless steel. An excellent bonding is also obtained in this case.

The temperature of the heat-treatment for bonding is such that the fusion-adhesion due to high Al layer with relatively low melting temperature of Al-alloy occurs, specifically, a temperature of 600°C or more. Note, since Fe diffuses from the core material into the coating layer simultaneously with the melting thereof, the coating layer instantaneously becomes an Fe-Al alloy, causing the melting point to rise and the layer to once solidify and to melt or nearly melt again at higher temperature, because Al content is very high in Fe-Al alloy layer at an early stage of diffusion and the melting temperature is not so high. The possible Al content of the Fe-Al alloy is 30 wt% at the highest after heat treatment, and the solidus of this alloy is 1300°C. The highest temperature for bonding treatment is, therefore, set at 1300°C.

Since the oxygen potential of the vacuum or inert gas-atmosphere becomes higher as the temperature becomes higher, the slight oxidation takes place even in the vacuum or inert gas atmosphere. Therefore, when the heat treatment for bonding is carried out at a high temperature, the formation of a thin  $Al_2O_3$  film occurs simultaneously with the fusion bonding, and the  $Al_2O_3$  film remains on the fusion-bonded honeycomb, and the like. The formation of an  $Al_2O_3$  film is, however, weak and the carrier blackens at around 700°C during the bonding treatment. In this case, the oxidizing treatment is carried out later at a temperature of from 850 to 1200°C.

Also, after brazing, it is evidently desirable to additionally form the  $Al_2O_2$  film by heating the brazed assembly in air. As described above, an outstanding feature of the present invention is that a stainless steel-foil having an Al-or Al-alloy-coating on the surface(s) is heat treated in vacuum or a reducing atmosphere, thereby simultaneously realizing a fusion-bonding between the contact parts, an improvement in the heat-resistance due to Al-diffusion from the coating layer to the core material, and the formation of the  $Al_2O_2$  film. If the bonding between the contacting parts of a honeycomb is unsatisfactory, a small amount of brazing material is supplementarily used to ensure the reliability of the bonding.

The present invention is further explained by way of examples.

55 Example I

30

	1					
5	-	Result ot Durability Bench Test	Abnormal oxidation occurs at brazed parts.	Normal	Normal.	Normal
15		reatme	: : 1140	<u>u</u>	я	in
20		Conditions for Bonding Treatment	Corrugated and flat sheets borded with brazing material (BNi5-grade). Treated at 1140°C in vacuum.	Heat treatment at 900°C in vacuum	Heat treatment at 600°C in vacuum	Heat treatment at 1000°C in hydrogen
<b>2</b> 5		ons fo	ted and with by rade). Jm.	eatmen	satment	atment
	Table 1	Conditic	Corrugated and bonded with br (BNi5-grade). in vacuum.	Heat tre vacuum	Heat tre vacuum	Heat tre hydrogen
30	Tat	/ethod	jation st and 1 70¢	mm osite 1. skness	ilzed 1885 1986 1988	hick
<b>3</b> 5		uring N	Corrugation ed sheet and ed, and eycomb 70\$	th 0.3 3 compo 50 µm 1 (thic	galvar stainle then r (thick	5 um thick both warm = 6/1
40		Honeycomb Manufacturing Method	Rolling to 50 µm. Corrugation forming. Corrugated sheet and flat sheet assembled, and coiled to form honeycomb $70\phi$ x 90 (mm) £	Al foil is clad with 0.3 mm thick stainless and composite foil then rolled to 50 µm. Stainless/Al = 10/1 (thickness ratio)	Al foil is hot-dip galvanized with 0.8 mm thick stainless and composite foil then rolled to 70 µm. Stainless/Al = 6/1 (thickness ratio)	to 50 µm. bonded to of SUS by s steel/Al ss ratio)
45		Hone	Roll form flat coil x 90	Al foi thick foil t Stainl ratio)	Al foi with 0 and co to 70 Stainl ratio)	Rolling that foils surfaces rolling. Stainless (thickness)
50	-	Substrate Material	15Cr-4Al	17Cr -0.3Ti -0.01C (XM)		25Cr-20N1 (SUS 310S)
55			Comparative Example	Inventive Examples		

5 .		Result of Durability Bench Test	<b>1</b>
			អុំ
15		Manufacturing Method Conditions for Bonding Treatment	Heat treatment at 1000°C in air. No bonding.
20		s for Bon	ument at ]
25	ontinued)	Condition	Heat treatm No bonding.
<b>30</b>	Table 1 (Continued)	Method	ł
35	LI	ıfacturing	um. 5 um ad to both 13 by warm 11/Al = 6/
40			ing to 50 oils bonde aces of St ing. nless stee ckness rat
45		e Honeycomb	i Roll f Surf surf roll Stai (thi
50		Substrate Material	arative '25Cr-20Ni Folling to 50 µm. 5 µm thick ample (5US Al foils bonded to both 310S) surfaces of SUS by warm rolling. Stainless steel/Al = 6/1 (thickness ratio)
55			arative ample

As shown in Table I, in one of the comparative examples, a rolled foil of I5Cr-4Al-steel was produced and shaped into a honeycomb, and the honeycomb was mounted in an outer cylinder made of AISI 304. Partial brazing was done for the flat and corrugated foils of the honeycomb, while brazing of the outer cylinder to the entire surface of the honeycomb periphery was carried out by using a BNi-5 grade brazing material, in vacuum at II40°C.

In one of the examples according to the present invention, Al foils (20  $\mu$ m thick) were bonded to both surfaces of a ferritic stainless-steel sheet (0.3 mm thick) containing I7Cr-0.3Ti-0.0IC, by warm rolling. Rolling was then carried out to obtain a composite foil of 50  $\mu$ m thickness. Subsequently, a honeycomb was manufactured and mounted in an outer cylinder made of AlSI 304. The brazing together of parts of the honeycomb components and of the outer cylinder and the honeycomb was carried out by a heat treatment at 900 °C in vacuum.

In another example according to the present invention, a 0.8 mm thick stainless-steel sheet having the same composition as that of the above example was subjected to hot-dip Al plating and then rolled to a thickness of 70  $\mu$ m. The honeycomb was shaped by the same method and then mounted in the outer cylinder. The heat treatment was carried out at 600°C in vacuum.

After the heat treatment, the above three honeycombs were subjected to a further treatment at  $900^{\circ}$ C for 7 hours in air to form the  $Al_2O_3$  film.

In the other example according to the present invention, 5  $\mu$ m thick AI foils were pressure-bonded on both surfaces of a 50  $\mu$ m thick foil of 25Cr-20Ni (AISI 3I0) by warm rolling, to obtain a clad material. The honeycomb was shaped by the same method and was mounted in the outer cylinder made of AISI 3I0. The heat treatment was carried out at 1000°C in vacuum. The same process was repeated except for a heat treatment at 1000°C in air, with regard to the other comparative example.

As a result, the bonding of honeycombs was attained by the heat treatment at a relatively low temperature, without the use of a brazing material, in the examples according to the present invention. The heat treatment in air did not result in a satisfactory bonding.

The brazed assemblies, including those of the comparative examples, were connected to and directly behind the exhaust manifold of engine having a displacement of  $1600\,\mathrm{cc}$ . The durability test was carried out over 30 hours at an output of 40 horse power and an engine rotation speed of 5000 rpm. The temperature of the exhaust gas was set in the range of from 940 to  $1000\,\mathrm{^oC}$  by delaying the ignition angle. Note,  $\gamma$ -Al<sub>2</sub>O<sub>3</sub> powder was not coated on the Al<sub>2</sub>O<sub>3</sub> film. As a result, abnormal oxidation occurred on portions of the brazed parts in the brazed assembly of the comparative example, but the brazed assemblies of the present invention remained totally normal and exhibited neither a peeling or the bonded parts nor deformation of the honeycombs.

### Example 2

20

25

Three cold-rolled strips of I5Cr-4.5Al steel, having a thickness of 0.4 mm were prepared. Al foils (40  $\mu$ m thick) were superimposed on both surfaces of one of the strips. The resultant three layers were bonded by cold rolling with a large reduction of 67% for one pass, and then rolled in the subsequent passes to a thickness of 50  $\mu$ m.

Al-alloy foils (Al-I0%Si-I%Mg) (40  $\mu$ m thick) were clad on both surfaces of another strip, and the resultant three layers were also rolled to obtain a composite foil of 50  $\mu$ m thickness.

The remaining steel strip was subjected to Al-alloy (Al-I0% Si) hot-dip plating. The amount of Al-alloy deposited per side was from 27 to 36  $\mu$ m. The thus-treated steel strip was then rolled to a thickness of 50  $\mu$ m.

As a result, three kinds of clad foils in total were obtained. These clad foils were used to form 5 honeycombs from each foil. Each honeycomb was pressed into an outer cylinder made of SUS 3IOS and heat-treated under the conditions shown in Table 2.

5 Protrusion at central part Conditions for Heat Treatment Bonding Condition Result of Bench Test No Abnormality No Abnormality No Abnormality No Abnormality No Abnormality 10 15 No Bonding 20 Bonded Bonded Bonded Bonded Bonded 25 Table 2-(1) Application of Ti powder Sealed in stainless foil at Application of Al powder. 1000°C x 1 hr in 58H,-Ar  $1000^{\circ}_{4}$ C x 1 hr in vacuum (10 Torr) 30 both ends of honeycomb 1200°C x 1 hr in air  $900^{\circ}$ C x 1 hr in dry H<sub>2</sub>  $1000^{\circ}$ C x 1 hr in air 1250°C x 1 hr in Ar 35 40 Structure Pure Al-Rolling Pure Al clad of -SUS-Foi.1 45 Comparative Invention 50

9

5 · · · · · · · · · · · · · · · · · · ·	Result of Bench Test	Protrusion at central part	No Abnormality .	No Abnormality	No Abnormality ,	No Abnormality	No Abnormalíty
20	Bonding Condition	No Bonding	Bonded	Bonded	Bonded	Bonded	Bonded
os Table 2-(2)	Conditions for Heat Treatment Bonding Condition Result of Bench Test	1000°C x 1 hr in air	$900^{\circ}$ C x 1 hr in dry H <sub>2</sub>	$1000^{\circ}$ C x 1 hr in $58H_2$ -Ar	Application of Al powder. 1250°C x l hr in Ar	Application of Ti powder Sealed in stainless foil at both ends of honeycomb 1200°C x 1 hr in air	1000°C x 1 hr in vacuum (10 Torr)
40	1	T000°C	× 2,006	1000°C	Applica 1250°C	Application Sealed : both end 1200°C ;	10004C 3
45	Foil Structure	Rolling clad of	AL ALIOY- -SUS-	AL alloy			
50		Comparative Rollin	Invention				

5		Conditions for Heat Treatment Bonding Condition Result of Bench Test	Protrusion at central part	No Abnormality	No Abnormality	No Abnormality	No Abnormality	No Abnormality
20		Bonding Condition	No Bonding	Bonded	Bonded	Bonded	Bonded	Bonded
<b>25</b>	Table 2-(3)	at Treatment	ir	ry H <sub>2</sub>	8H <sub>2</sub> -Ar powder.	Ar	powder ss foil at ycomb air	acuum
<i>30</i>		onditions for He	650°C x l hr in air	$650^{\circ}$ C x 1 hr in dry H <sub>2</sub>	800°C x 1 hr in 58H2-Ar Application of Al powder.	1000°C x l hr in Ar	Application of Ti powder Sealed in stainless foil at both ends of honeycomb 1000°C x 1 hr in air	650°C x l hr in vacuum (10 <sup>-4</sup> Torr)
40		Foil Structure	Rolling 6 cladding,	1	ŀ	-	<b>&amp;</b> & & C	9
50			Comparative	Invention				

After the heat treatment, the bonding conditions of the brazed assemblies were confirmed by the same bench test by an engine as in Example I.

#### 0 243 702

As a result it was recognized that, in the direct heat treatment in air (Comparative Examples) bonding of the honeycombs was not realized, and that an abnormality occurred, i.e., the central part of the honeycombs protruded in the direction of flow of the exhaust gas, as a result of the bench test. Contrary to this in the heat treatments in dry H<sub>2</sub>, 5%H<sub>2</sub>-Ar, vacuum, Ar gas (after application of Al powder), and in air (after application of Ti powder and sealing by stainless steel foils at both ends of the honeycomb), the bonding of honeycombs was realized and the abnormality appearing in the Comparative Examples did not occur.

As described hereinafter, the present invention makes it possible to carry out bonding a honeycomb for the exhaust gas purification device of an automobile, by an extremely simple method, in place of the conventional brazing method. Further, in this method, the honeycomb is given a very high heat resistance. Accordingly, it further contributes to a cost reduction and an enhancement of the characteristics of such a catalyst converter.

#### 75 Claims

- I. A method for producing a base, on which a catalyst carrier for purifying exhaust gas of an automobile is deposited, wherein a stainless-steel sheet foil is shaped into a honeycomb, and the peripheral surface of the honeycomb is bonded to an outer cylinder, characterized by rolling the stainless-steel sheet having a layer of Al or Al-alloy on one or both surfaces of the stainless-steel sheet to form a composite foil, producing the honeycomb structure by using the composite foil, mounting the honeycomb structure in the outer cylinder of heat-resistant steel and heat-treating the honeycomb and outer cylinder in vacuum or a reducing atmosphere at a temperature of from 600 to 1300°C.
- 2. A method according to claim I, wherein the stainless-steel sheet to be provided with the AI or AI-alloy layer thereon, is an AI-containing stainless steel strip.
  - 3. A method according to claim I or 2, wherein the layer of AI or AI-alloy is a foil layer.
- 4. A method according to claim I or 2, wherein the layer of Al or Al-alloy is a hot-dip plating layer of Al or Al-alloy.
- 5. A method according to claim I, wherein the reducing atmosphere is formed by a reducing agent deposited on the surfaces of honeycomb.
  - 6. A method according to claim I, wherein the Al-alloy contains one or more of Mn, Mg. Si, and Cu.

35

40

45

50

# **EUROPEAN SEARCH REPORT**

Application number

EP 87 10 4765

DOCUMENTS CONSIDERED TO BE RELEVANT						
Category	Citation of document with of relevi	n indication, where appropriate, ant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.4)		
A	GB-A-2 081 747 ( * Figure 2; colum column 4, line 11	n 2, line 10 -	1-3	F 01 N 3/28 B 01 D 53/36		
A	FR-A-2 321 347 (ENERGY AUTHORITY)					
A	 EP-A-0 159 468 (	INTERATOM)				
A	FR-A-2 470 246 (	BREMSHEY AG)				
A	DE-A-2 319 663 ( AG)					
Control of the Contro				TECHNICAL FIELDS SEARCHED (Int. Cl.4)		
				F 01 N 3/00 B 01 D 53/00		
	·					
	The present search report has b	een drawn up for all claims				
	Place of search THE HAGUE	Date of completion of the search 20-07-1987	BOGA	Examiner ERTS M.L.M.		
Y:pd	CATEGORY OF CITED DOCL articularly relevant if taken alone articularly relevant if combined wocument of the same category schnological background on-written disclosure termediate document	E : earlier pat after the fi ith another . D : document L : document	ent document ling date cited in the a cited for othe f the same pat	erlying the invention t, but published on, or pplication er reasons tent family, corresponding		

THIS PAGE BLANK (USPTO)